CLEAN BREATH TOOTH- precipitated calcium carbonate, dibasic calcium phosphate, aminocaproic acid, aluminium chlorohydroxy allantoinate paste, dentifrice
K.Boeun Pharmaceutical Co.,Ltd.

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

Precipitated Calcium Carbonate, Dibasic Calcium Phosphate, Aminocaproic Acid, Aluminium Chlorohydroxy Allantoinate

Water, D-Sorbitol Solution, Concentrated Glycerin, Silicon Dioxide, Sodium Cocoyl Glutamate, Polyethylene Glycol 1500, Xanthangum, Mentha Oil, Enzymatically Modified Stevia, Titanium Oxide, Sodium Chloride, Propolis Extract, l-Menthol, Xylitol, Ascorbic Acid, Green Tea Extract, Aloe Extract, Tocopherol Acetate

For dental care

Keep out of reach of children

Apply an appropriate amount to your toothbrush and brush your teeth by brushing.

Warnings

Keep out of reach of children

■ If more than used for rinsing is accidentally swallowed, get medical help or contact a Poison Control Center right away.

Other Information

■ Store in an airtight container at room temperature

■ Date of use : 36 months from the date of manufacture

For dental use only

CLEAN BREATH TOOTH

precipitated calcium carbonate, dibasic calcium phosphate, aminocaproic acid, aluminium chlorohydroxy allantoinate paste, dentifrice

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:74724-0024
Route of Administration	DENTAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
CALCIUM PHOSPHATE, DIBASIC, ANHYDROUS (UNII: L11K75P92J) (CALCIUM PHOSPHATE, DIBASIC, ANHYDROUS - UNII:L11K75P92J)	CALCIUM PHOSPHATE, DIBASIC, ANHYDROUS	0.07 g in 100 g

CALCIUM CARBONATE (UNII: H0G9379FGK) (CALCIUM CATION - UNII:2M83C4R6ZB)	CALCIUM CARBONATE	36 g in 100 g
ALCLOXA (UNII: 18B8O9DQA2) (ALCLOXA - UNII:18B8O9DQA2)	ALCLOXA	0.06 g in 100 g
AMINOCAPROIC ACID (UNII: U6F3787206) (AMINOCAPROIC ACID - UNII:U6F3787206)	AMINOCAPROIC ACID	0.06 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
XYLITOL (UNII: VCQ006KQ1E)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:74724-0024-1	30 g in 1 TUBE; Type 0: Not a Combination Product	05/01/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		05/01/2021	

Labeler - K.Boeun Pharmaceutical Co.,Ltd. (695674074)

Registrant - K.Boeun Pharmaceutical Co.,Ltd. (695674074)

Establishment

Name	Address	ID/FEI	Business Operations
K.Boeun Pharmaceutical Co.,Ltd.		695674074	manufacture(74724-0024)

Revised: 5/2021 K.Boeun Pharmaceutical Co.,Ltd.